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<p>(54) Title: NEW SULFONE ESTER ANALOGUES OF iso-CLOZAPINE AND RELATED STRUCTURES: ATYPICAL NEUROLEPTICS</p> <p>(57) Abstract</p> <p>A compound of formula (I), or pharmaceutically acceptable acid addition salts thereof, wherein R₁ is H, (C₁-C₈) alkyl or haloalkyl or hydroxyalkyl, alkenyl, alkynyl, cyclopropylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl; R₂ is H, (C₁-C₈) alkyl, alkenyl, alkynyl, cyclopropylalkyl or (C₁-C₈) haloalkyl, hydroxyalkyl, hydroxyalkyloxyalkyl or 1-(alkyl-2-imidazolidinone); X is NH, NR₁, O, S, SO₂. The compounds of this invention possess affinity to one or several receptor systems, e.g. DA (D1-D4), α₁, muscarinic (M1-M4) and 5-HT (5-HT2A, 5-HT2C and 5-HT7). The central nervous system disorders to be treated with the compounds of the present invention include psychoses-schizophrenia, autism, Tourette's syndrome, restless legs, Huntington's chorea, motion sickness, nausea, vomiting and severe anxiety.</p>			
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NEW SULFONE ESTER ANALOGUES OF iso-CLOZAPINE AND
RELATED STRUCTURES: ATYPICAL NEUROLEPTICS.

Field of the Invention

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The present invention is directed toward new sulfone ester analogues of iso-Clozapine [RN 1977-08-8] and their pharmaceutically acceptable salts, to processes for preparing such compounds, pharmaceutical preparations of such compounds and the use of such compounds in manufacture of a pharmaceutical preparation. Pharmaceutical preparations of these compounds are useful for central nervous system disorders, such as for a therapeutic effect on one or several of the following receptor systems: dopamine (DA; D1-D4), 5-HT2A, 5-HT2C, α 1 and muscarinic (M1-M4) in mammals.

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In psychotic/schizofrenic patients evidence indicates that the neurotransmission in the central nervous system (CNS) may be disturbed. These disturbances seem to involve the neurotransmitters dopamine (DA, D1-D4 receptors), noradrenaline (NA, α 1 receptors), acetylcholine (ACh, M1-M4 receptors) and 5-hydroxytryptamine (5-HT, 5-HT2A and 5-HT2C receptors).¹ The drugs most frequently used in the treatment of psychoses are considered to act by altering the neurotransmission of one or several of these receptor systems. The mechanism of action for typical/classical anti-psychotic drugs used to treat psychosis is generally believed to be through blockage of DA D2 receptors.^{2, 3, 4} The atypical anti-psychotic drug Clozapine [RN 54241-01-9] is an efficient anti-psychotic agent⁵ and binds with high affinity to DA D1 and D4 receptors and with moderate affinity to DA D2 and D3 receptors.⁶ In addition, Clozapine binds with high affinity to α 1 receptors, muscarinic M1-M4 receptors and 5-HT2A and 5-HT2C receptors.⁶ Clozapine has a low propensity to induce extrapyramidal side-effects in humans and catalepsy in rats. In addition, Clozapine does not increase the prolactin levels, something which the classical neuroleptics (e. g. haloperidol and chlorpromazine) potently do. However, the use of Clozapine is hampered by rare incidences of a serious side-effect (blood toxicity, agranulocytosis).⁶ Expensive, continuous blood monitoring makes the use of Clozapine tedious and expensive and the use of Clozapine is therefore limited to psychotic patients not responding to the first and second choice of drug therapy.⁷

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Several Clozapine analogues have been evaluated but, to date, no marketed compound with a Clozapine-like structure and Clozapine-like pharmacological and clinical profile, i. e.

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an analogue that is also free from side-effects, has been found.⁷

A very interesting and, for the present invention, a highly relevant fact is that the 2-Cl isomer of Clozapine (iso-Clozapine, [RN 1977-08-8]) has been classified as a typical neuroleptic agent.⁸ We surprisingly found that the iso-Clozapine sulfone analogues of the present invention display atypical neuroleptic properties.

Information Disclosure Statement.

10 The following documents could be important in the examination of this application:

iso-Clozapine references: 9, 10, 11

Loxapine references: 12, 13, 14, 15, 16, 17, 18, 19, 20, 21

15 Clothiapine references: 15, 16, 19, 21, 22, 23, 24, 25, 26

Miscellaneous references over sulfonates: 27, 28, 29, 30, 31

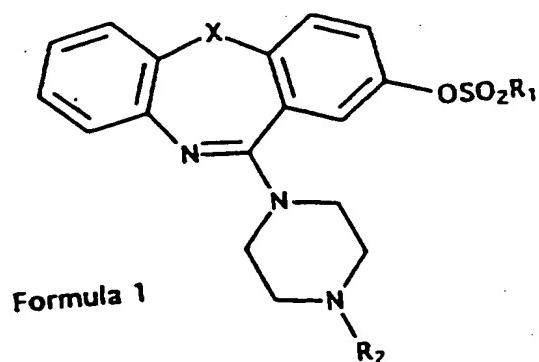
20 Sulfonate patent application reference: 32

Summary of the Invention

This invention is related to novel iso-Clozapine, Loxapine and Clothiapine analogues of

25 Formula 1:

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or pharmaceutically acceptable acid addition salts thereof, wherein R₁ is H, (C₁-C₈) alkyl or haloalkyl (e. g. CF₃ (triflates) and CF₂CF₃) or hydroxyalkyl, alkenyl, alkynyl,

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cyclopropylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl; R₂ is H, (C₁-C₈) alkyl, alkenyl, alkynyl, cyclopropylalkyl or (C₁-C₈) haloalkyl (e. g. CH₂CF₃, CH₂CH₂CF₃, CH₂CH₂F, CH₂CH₂CH₂F), hydroxyalkyl, hydroxyalkyloxyalkyl (e. g. CH₂CH₂OCH₂CH₂OH) or 1-(alkyl-2-imidazolidinone) (e. g. 1-(CH₂CH₂-2-imidazolidinone) and 1-(CH₂CH₂CH₂-2-imidazolidinone)); X is NH, NR₁, O, S, SO, SO₂.

The compounds of this invention possess affinity to one or several receptor systems, e. g. DA (D1-D4), α 1, muscarinic (M1-M4) and 5-HT (5-HT2A, 5-HT2C and 5-HT7). The central nervous system disorders to be treated with the compounds of the present invention include psychoses/schizophrenia, autism, Tourette's syndrome, restless legs, Huntington's chorea, motion sickness, nausea, vomiting and severe anxiety.

In a preferred embodiment, the invention is directed to compounds of Formula 1 wherein R₁ is haloalkyl R₂ is H or CH₃ and X is NH, O or S. A more preferred embodiment are compounds of Formula 1 wherein R₁ is CF₃, R₂ is H or CH₃ and X is NH, O or S. An even more preferred embodiment are compounds of Formula 1 wherein R₁ is CF₃, R₂ is CH₃ and X is NH, O or S. The most preferred compound is the compound of Formula 1 wherein R₁ is CF₃, R₂ is CH₃ and X is NH.

An object of the invention is to provide compounds for therapeutic use, especially compounds having a therapeutic activity in the central nervous system. Another object is to provide compounds having an effect on receptors (DA (D1-D4), α 1, muscarinic (M1-M4) and 5-HT (5-HT2A, 5-HT2C and 5-HT7)) in mammals including man.

Processes for preparation of these compounds, their pharmaceutical use and pharmaceutical preparations employing such compounds constitute further aspects of the invention.

Detailed Description of the Invention

In appropriate situations, the proper stereochemistry is represented in the structural schemes.

As used herein the term (C_n-C_m) is inclusive such that a compound of (C₁-C₈) would include compounds of one to 8 carbons and their isomeric forms. The various carbon moieties are defined as follows: alkyl refers to an aliphatic hydrocarbon radical and

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includes branched or unbranched forms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, n-pentyl, isopentyl, neo-pentyl, n-hexyl, isohexyl, n-heptyl, isoheptyl, and n-octyl.

- 5 Alkenyl refers to a radical of an aliphatic unsaturated hydrocarbon having a double bond and includes both branched and unbranched forms such as ethenyl, 1-methyl-1-ethenyl, 1-propenyl, 2-propenyl, 1-butenyl, 2-butenyl, 3-butenyl, 2-methyl-1-butenyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-methyl-4-pentenyl, 3-methyl-1-pentenyl, 3-methyl-2-pentenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 1-methyl-4-hexenyl, 3-methyl-1-hexenyl, 3-methyl-2-hexenyl, 1-heptenyl, 2-heptenyl, 3-heptenyl, 4-heptenyl, 1-methyl-4-heptenyl, 3-methyl-1-heptenyl, 3-methyl-1-heptenyl, 3-methyl-2-heptenyl, 1-octenyl, 2-octenyl or 3-octenyl.

10 Cycloalkyl refers to a radical of a saturated cyclic hydrocarbon such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, or cyclooctyl. "Halogen" means fluorine, chlorine, bromine or iodine, preferably fluorine.

- 15 It will be apparent to those skilled in the art that compounds of this invention may contain chiral centers (e. g. R₁ and/or R₂). The scope of this invention includes all enantiomeric or diastereomeric forms of Formula 1 compounds, either in pure form or as mixtures of enantiomers or diastereomers. The compounds of Formula 1 can contain one asymmetric carbon atom in the ring system (when X=SO). The therapeutic properties of the compounds may to a greater or lesser degree depend on the stereochemistry of a particular compound. Pure enantiomers, as well as enantiomeric or diastereomeric mixtures, are within the scope of the invention.
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- 25 Both organic and inorganic acids can be employed to form non-toxic pharmaceutically acceptable acid addition salts of the compounds of this invention. Illustrative acids are sulfuric, nitric, phosphoric, hydrochloric, citric, acetic, lactic, tartaric, pamoic, methanesulfonic, ethanesulfonic, sulfamic, succinic, cyclohexylsulfamic, fumaric, maleic and benzoic acid. These salts are readily prepared by methods known in the art.

- 30 The compounds of this invention may be obtained by one of the following methods described below, as outlined in the appropriate charts.

- 35 In clinical practice the compounds of the present invention will normally be administered orally, rectally or by injection, in the form of pharmaceutical preparations comprising the active ingredient either as a free base or as a pharmaceutically acceptable non-toxic, acid addition salt, such as the hydrochloride, lactate, acetate, methanesulfonate, sulfamate

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salt, in association with a pharmaceutically acceptable carrier. The use and administration to a patient to be treated in the clinic would be readily apparent to a person of ordinary skill in the art.

5 In therapeutical treatment the suitable daily doses of the compounds of the invention are from about 0.1 mg to 2000 mg for oral application, preferably 0.5-500 mg, and 0.05 mg to about 100 mg for parenteral application, preferably 0.05-50 mg daily doses. The daily dose will preferably be administrated in individual dosages 1-4 times daily and the dosage amounts are based on an individual having a weight of 70 kg.

10 The compounds of Formula 1 of this invention, wherein R₁ is H, (C₁-C₈) alkyl or haloalkyl (e. g. CF₃ (triflates) and CF₂CF₃) or hydroxyalkyl, alkenyl, alkynyl, cyclopropylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl; R₂ is H, (C₁-C₈) alkyl, alkenyl, alkynyl, cyclopropylalkyl or (C₁-C₈) haloalkyl (e. g. CH₂CF₃, CH₂CH₂CF₃, CH₂CH₂F,

15 CH₂CH₂CH₂F), hydroxyalkyl, hydroxyalkyloxyalkyl (e. g. CH₂CH₂OCH₂CH₂OH) or 1-(alkyl-2-imidazolidinone) (e. g. 1-(CH₂CH₂-2-imidazolidinone) and 1-(CH₂CH₂CH₂-2-imidazolidinone)); X is NH, NR₁, O, S, SO, SO₂, display binding affinity at the receptor systems: DA (D1-D4), α1, muscarinic (M1-M4) and 5-HT (5-HT_{2A}, 5-HT_{2C} and 5-HT₇) in mammals including man. These compounds are particularly effective atypical anti-psychotic (anti-schizophrenic) agents. Other uses for these compounds include autism, Tourette's syndrome, restless legs, Huntington's chorea, motion sickness, nausea, vomiting and severe anxiety.

The compounds of this invention also have high oral potency and a long duration of action.

The utility of the compounds of this invention to treat central nervous system disorders can be shown in behavioral and biochemical activity in rats.

30 Effect on DA release in rats as measured by microdialysis in freely moving animals.

The biochemical effects of the compounds of this invention can be monitored with the microdialysis technique in freely moving animals (see under Experimental Section below).

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Monitoring potential catalepsy in rats.

The propensity of the compounds of this invention to induce catalepsy was measured in rats, who were placed temporarily on a vertical grid (see under Experimental Section 5 below).

Monitoring of potential inhibition of locomotor activity in rats treated with Apomorphine or Amphetamine.

10 The ability of the compounds of this invention to reduce the Apomorphine or Amphetamine induced increase in locomotor activity was measured in rats with motility meters (see under Experimental Section below).

Experimental Section

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Syntheses

Starting materials (the corresponding phenols) for the claimed target compounds may b obtained by the methods known in the art.

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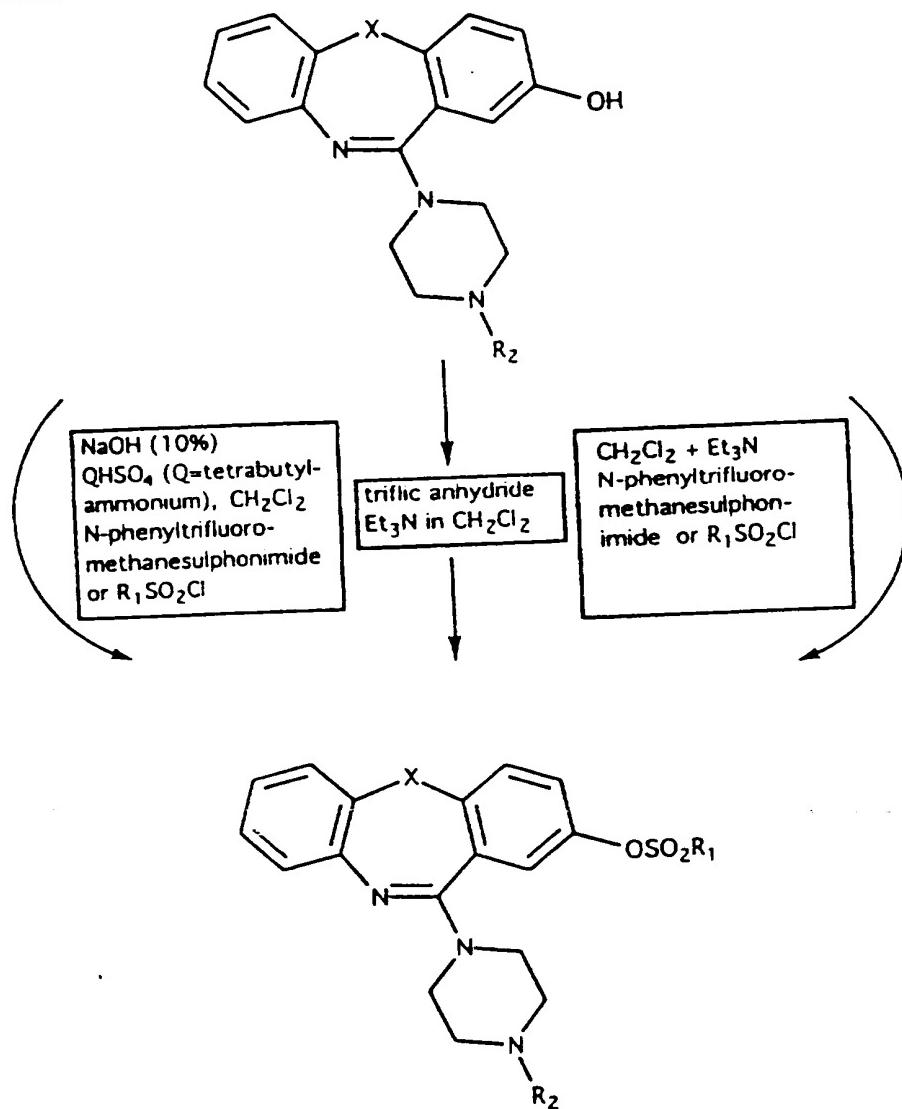
¹H and ¹³C NMR spectra were recorded at 200 and 50.3 MHz, respectively, on a Varian Gemini 200 spectrometer. CDCl₃ was employed as the solvent unless otherwise stated. Chemical shifts are given in units (ppm) and relative to TMS or deuterated solvent. IR spectra were obtained on a ATI-Mattson spectrometer. Elemental analyses were performed in the Micro Analytical Department of the University of Groningen or at other approved laboratories. The chemical ionization (CI) mass spectra were obtained on a Finnegan 3300 system or a UNICAM Automass 150 GC/MS system, equipped with a UNICAM GC, 610 Series. Melting points were determined on a Electrothermal digital melting point apparatus and are uncorrected. Specific optical rotations were measured in methanol (c = 1.0) at 21 °C on a Perkin Elmer 241 polarimeter.

All methoxylated starting compounds were prepared according to the literature procedures.^{10, 18, 21, 33, 34} Chemicals used were commercially available (Aldrich and/or Sigma) and were used without further purification.

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Parts of the synthetic methods, not necessarily the preferred route, for forming the sulfone esters from the corresponding phenols are exemplified in Scheme 1:

Scheme 1

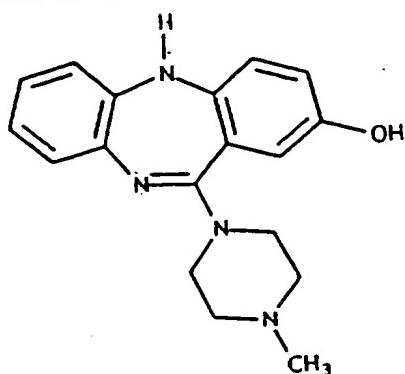


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Suitable starting materials, either the phenols or the OMe-analogues, which can be dealkylated to the corresponding phenols with e. g. AlCl₃ in ethylmercaptane at room temperature, for this invention are known:

- 5 2-Hydroxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine; 2-hydroxy-8-dechloroclozapine; RN [156632-07-4]

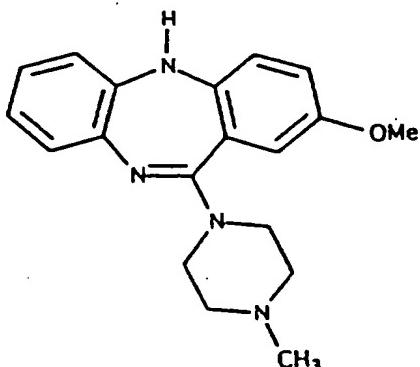
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- 2-Methoxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine; 2-methoxy-8-dechloroclozapine; RN [95316-97-5]

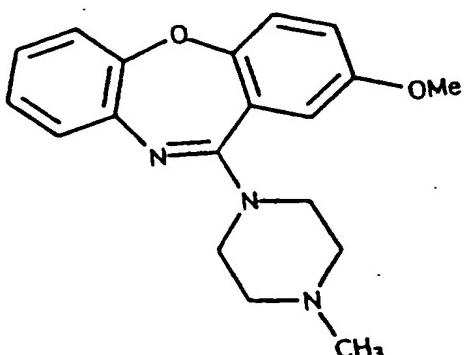
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- 2-Methoxy-11-(4-methyl-1-piperazinyl)-dibenzo[b,f][1,4]oxazepine; RN [13745-79-4]

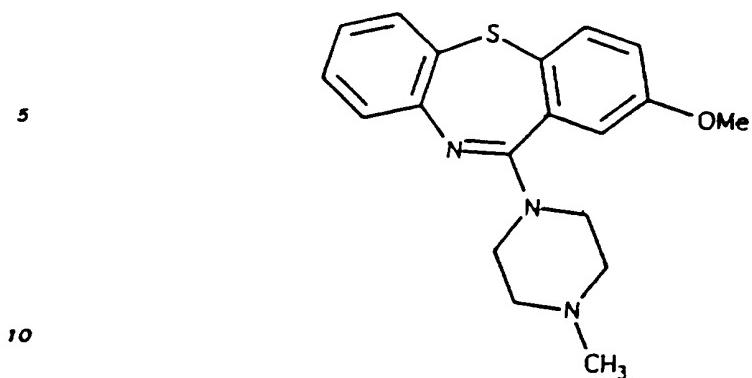
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2-Methoxy-11-(4-methyl-1-piperazinyl)-dibenzo[b,f][1,4]thiazepine; RN [13745-79-4]



The following intermediates, useful for the preparation of compounds of the present invention are known:

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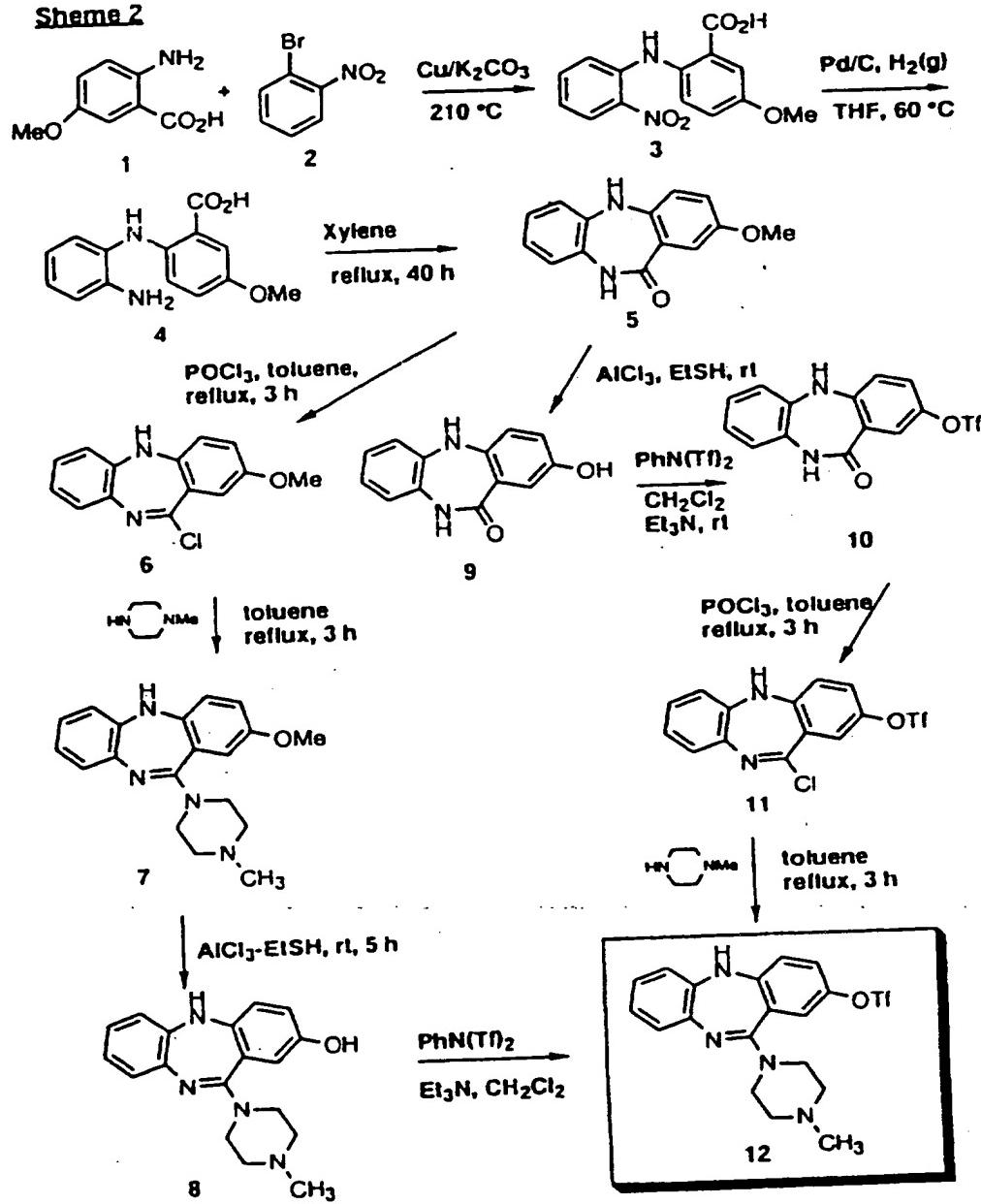
2-Methoxy-dibenzo[b,f][1,4]oxazepin-11(10H)-one RN [60287-33-4]³⁵

2-Hydroxy-dibenzo[b,f][1,4]oxazepin-11(10H)-one RN [60287-08-3]³⁵

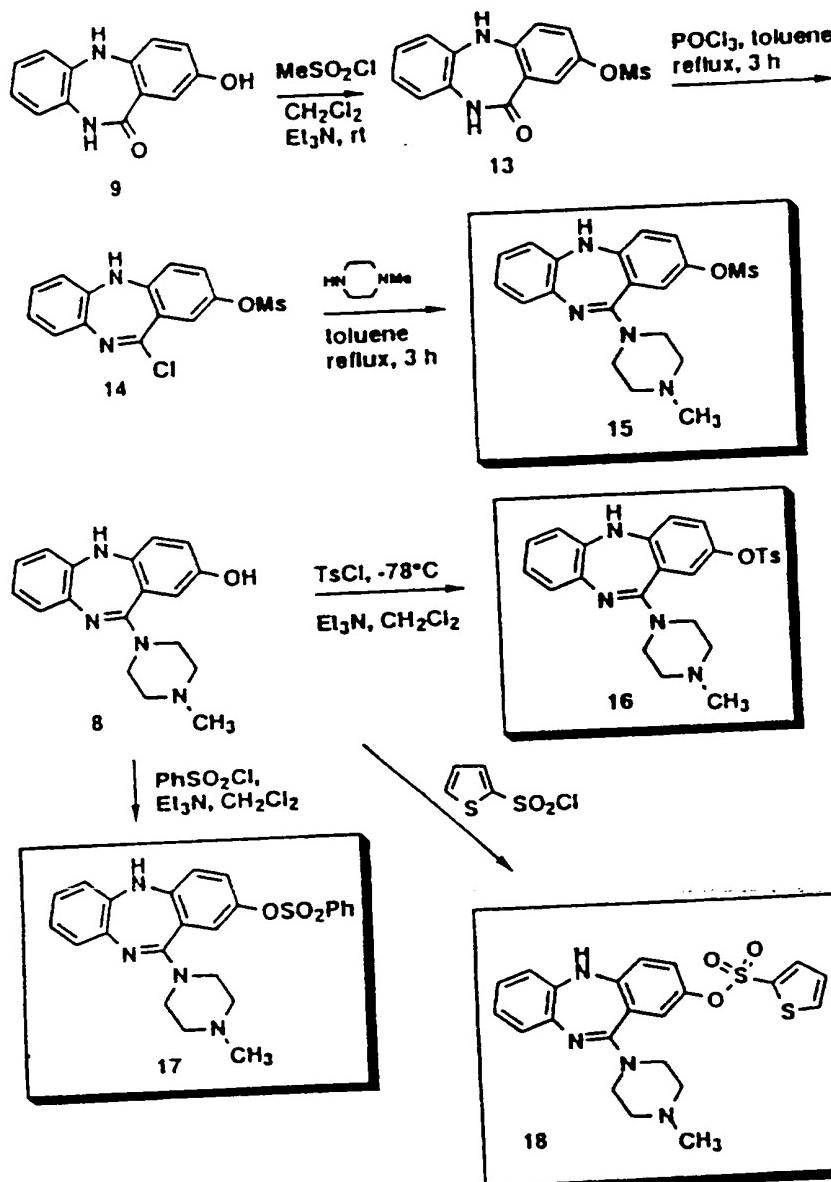
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2-Methoxy-dibenzo[b,f][1,4]thiazepin-11(10H)-one RN [3158-77-8]^{36, 37}

- 10 -

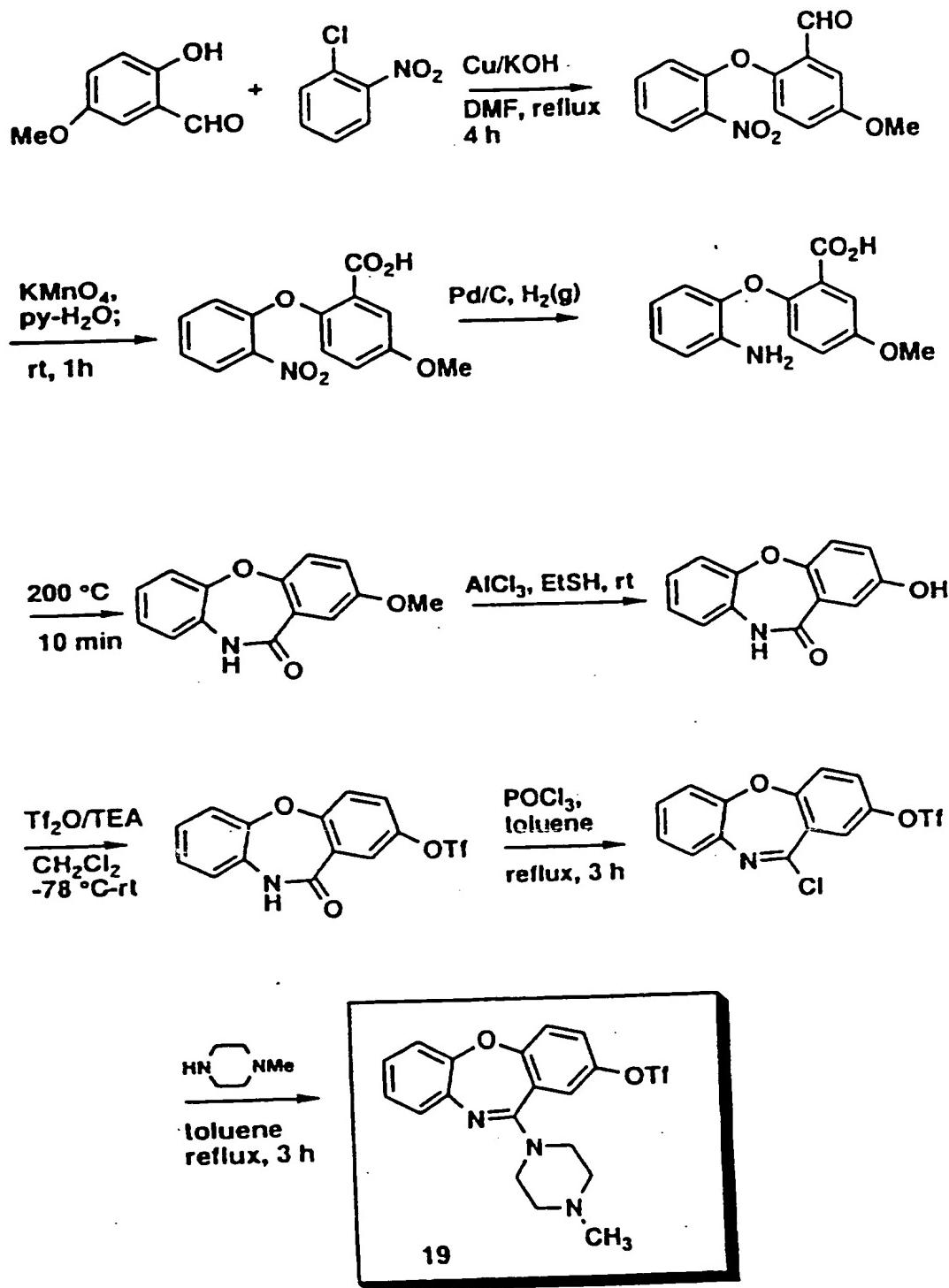
Scheme 2

Scheme 3

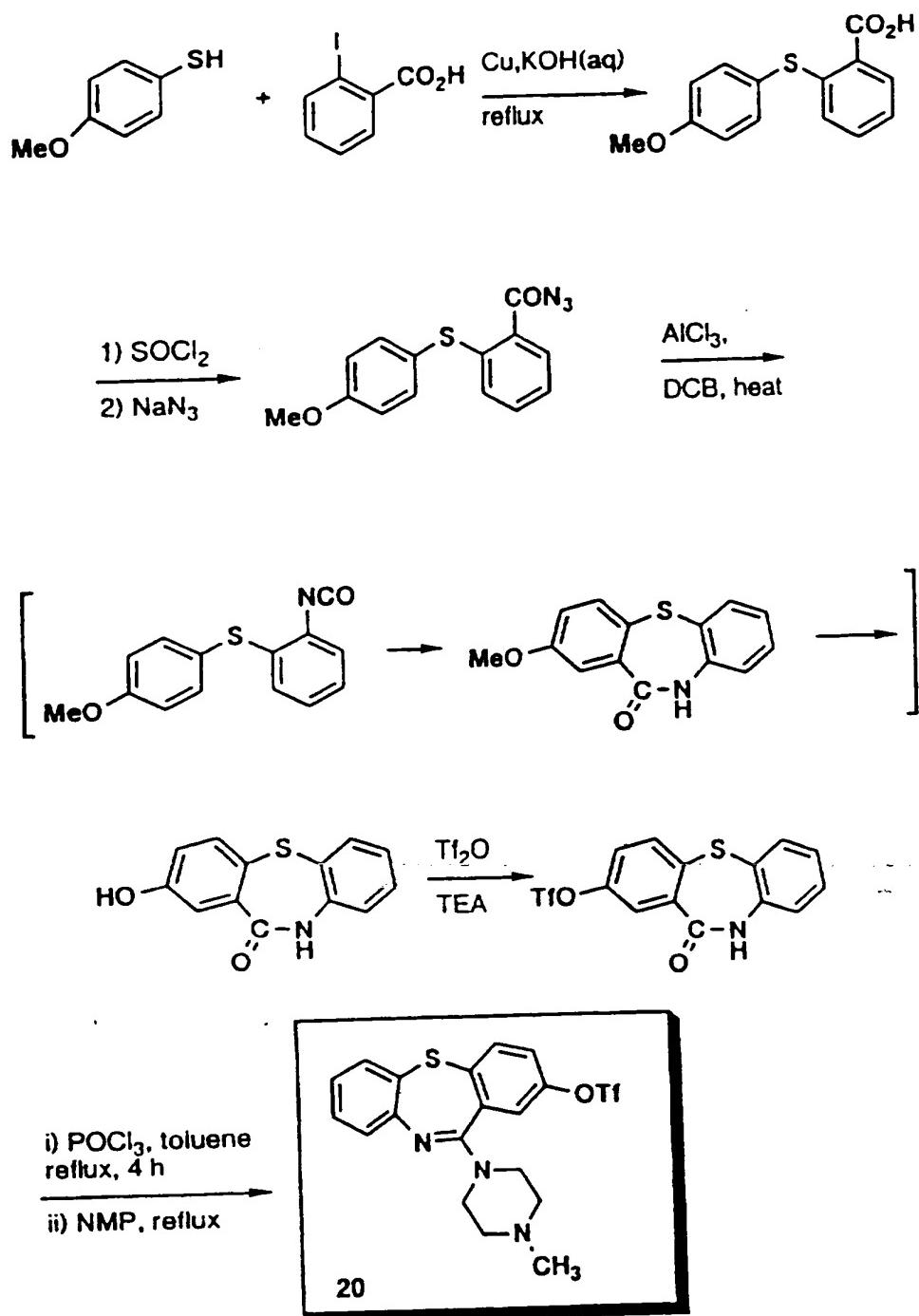


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Scheme 4



Scheme 5



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Synthesis of 2-Trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (12).

2-(2-Nitrophenyl)amino-5-methoxybenzoic acid (3). A mixture of 5-methoxy-2-aminobenzoic acid (1, 6.68 g, 40 mmol), 2-bromo-nitrobenzene (2, 8.08 g, 40 mmol) and 10 mL of amyl alcohol was heated to 80 °C with stirring. To the homogeneous solution was immediately added copper powder (60 mg) and K₂CO₃ (5.7 g). The reaction temperature was increased in order to distill off the water generated in the reaction and the amyl alcohol. The reaction mixture was then kept at 210 °C for 3 h. After cooling, the reaction mixture was quenched with 200 mL of water and 100 mL of 4 N aq HCl at room temperature. The precipitate was removed by filtration and the solution was extracted with ethyl acetate. The organic layers were combined and concentrated. The residue was purified by flash chromatography (SiO₂, ethyl acetate as eluent) to afford 6.8 g (59 %) of the title compound.

2-(2-Aminophenyl)amino-5-methoxybenzoic acid (4). Compound 3 [2-(2-nitrophenyl)amino-5-methoxybenzoic acid] (1.8 g) in THF (50 mL) was hydrogenated with a catalytic amount of palladium on carbon in a Parr-shaker apparatus under 4.5 atm H₂ (g) at 60 °C for 4 h to afford 1.4 g (88 %) of the title compound.

2-Methoxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]diazepine (5). Intermediate 4 [2-(2-aminophenyl)amino-5-methoxybenzoic acid] (3.0 g) in 100 mL of xylene was refluxed for 40 h. After evaporation the residue was purified by flash chromatography (SiO₂, hexane and ethyl acetate, 4:1 then 1:1, as eluents) to afford 2.2 g (79 %) of the title compound.

MS (EI) shows M⁺ at m/e = 240.

2-Methoxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]-diazepine (7, RN [95316-97-5]). Intermediate 5 (200 mg, 0.83 mmol), phosphorus oxychloride (POCl₃) (3 mL), toluene (5 mL) and N,N-dimethylaniline (0.5 mL) were combined and heated to reflux for 3 h. The solvents of the mixture were evaporated under vacuum to afford the iminochloride intermediate 6, which was used in the next-step reaction without further purification.

Above iminochloride 6 in 3 mL of toluene and 2 mL of N-methylpiperazine were refluxed for 3 h. After evaporation the residue was taken up with chloroform, washed with 2N aq NaOH and purified by flash chromatography (SiO₂, ethyl acetate then THF as eluents) to afford 214 mg (80 %) of the title compound. MS (EI) shows M⁺ at m/e = 322.

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2-Hydroxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]-diazepine (8, RN [156632-07-4]). 2-Methoxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (7) (200 mg, 0.62 mmol) in 3 mL of EtSH was treated with AlCl₃ (1.5 g) with stirring at room temperature for 4 h. The mixture was quenched with 30 mL of ice-water. The pH value of the solution was adjusted to pH 8 with 2N aq NaOH. The solution was extracted with chloroform (4 x 30 mL). The organic layers were combined, dried over MgSO₄, filtered, and the solvents were evaporated. The light yellow solid residue was recrystallized from ethyl acetate to afford 140 mg (73 %) of the title compound as fine crystals: mp 266 °C, with all of the expected spectra including IR, ¹H NMR, ¹³C NMR. MS (EI) shows M⁺ at m/e = 308.

2-Hydroxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]diazepine (9). 2-Methoxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]diazepine (5) (600 mg, 2.5 mmol) in 4 mL of EtSH was treated with AlCl₃ (1.6 g) with stirring at room temperature for 4 h. The mixture was quenched with 20 mL of ice-water and then 10 mL of 4N aq HCl. The solution was extracted with chloroform (2 x 20 mL) and ethylacetate (2 x 20 mL). The combined organic layers were dried over MgSO₄, filtered, and the solvents were evaporated. The solid residue was recrystallized from ethyl acetate to afford 505 mg (88 %) of the title compound as fine crystals: mp 280 °C, with all the expected spectra including IR, ¹H NMR, ¹³C NMR. MS (EI) shows M⁺ at m/e = 226.

2-Trifluoromethanesulfonyloxy-5,10-dihydro-11-oxo-dibenzo-[b,e][1,4]diazepine (10). 2-Hydroxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]-diazepine (9) (400 mg, 1.77 mmol) in 10 mL of dioxane was treated with N-phenyl-trifluoromethanesulfonimide (800 mg) in the presence of 2 mL of triethylamine with stirring at room temperature overnight. After evaporation of the solvents, the residue was purified by flash chromatography (SiO₂, hexane and ethyl acetate, 9:1 then 4:1, as eluents). Recrystallization from hexane and ethyl acetate (10:1) afforded 380 mg (60 %) of the title compound as light yellow crystals. MS (EI) shows M⁺ at m/e = 358.

2-Trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (12). Method A. 2-Hydroxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (8) (180 mg, 0.58 mmol) in 10 mL of CH₂Cl₂ was treated with N-phenyl-trifluoromethanesulfonimide (300 mg, 0.83 mmol) in the presence of 1 mL of triethylamine with stirring at room temperature overnight. After evaporation the residue was purified by flash chromatography (SiO₂; hexane and ethyl acetate, 1:1; then ethyl

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acetate as eluents). Recrystallization from hexane and ethyl acetate (15:1) afforded 200 mg (78 %) of the title compound as fine crystals: mp 160 °C, with the expected spectra data including IR, ¹H NMR, ¹³C NMR. MS (EI) shows M⁺ at m/e = 440.

5 **Method B.** 2-Trifluoromethanesulfonyloxy-5,10-dihydro-11-oxo-dibenzo-[b,e][1,4]diazepine (10) (260 mg, 0.72 mmol), phosphorus oxychloride (POCl₃) (4 mL), toluene (10 mL) and N,N-dimethylaniline (0.5 mL) were combined and heated to reflux for 3 h. The mixture was evaporated under vacuum to afford the corresponding iminochloride intermediate 11, which was used in the next-step reaction without further purification.

10 The iminochloride 11 in 10 mL of toluene was treated with 3 mL of N-methylpiperazine under refluxing for 3 h. After evaporation of the solvents, the residue was taken up in chloroform, washed with 2N aq NaOH and purified by flash chromatography and recrystallized, as described in Method A above, to afford 230 mg (73 %) of the title 15 compound 12.

2-Methanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo-[b,e][1,4]diazepine (15).

20 **2-Methanesulfonyl-5,10-dihydro-11-oxo-dibenzo(b,e)[1,4]diazepine (13).** 2-Hydroxy-5,10-dihydro-11-oxo-dibenzo(b,e)[1,4]diazepine (9) (50 mg, 0.22 mmol) in 6 mL of dichloromethane was treated with methanesulfonylchloride (34 mg, 0.30 mmol) in the presence of 0.2 mL of triethylamine at -78 °C. The mixture was stirred for 1 h, in order to allow the reaction temperature to reach room temperature, quenched with water and extracted with chloroform. Workup afforded 50 mg (76 %) of the title compound. MS (EI) 25 shows M⁺ at m/e = 304.

30 **2-Methanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (15).** 2-Methanesulfonyloxy-5,10-dihydro-11-oxo-dibenzo(b,e)[1,4]diazepine (13) (50 mg, 0.16 mmol), phosphorus oxychloride (POCl₃) (2 mL), toluene (3 mL) and N,N-dimethylaniline (0.1 mL) were combined and heated to reflux for 3 h. The mixture was evaporated under vacuum to afford the iminochloride intermediate 14, which was used in the next reaction step without further purification.

35 This iminochloride 14 in 3 mL of toluene was treated with 1 mL of N-methylpiperazine under refluxing for 3 h. Similar workup as described for the preparation of 12 afforded 32 mg (50 %) of the title compound as fine crystals: mp 173-177 °C. MS (EI) shows M⁺ at m/e = 386.

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Preparations of 2-(4-methylphenylsulfonyloxy)-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepin (16), 2-phenylsulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (17) and 2-(2-thienylsulfonyloxy)-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (18) were performed by the same procedure as for the preparation of 2-trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (12) (Method A) from 2-hydroxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo[b,e][1,4]diazepine (8), however, using the sulfonation procedure with appropriate sulfonylchloride, as described for the preparation of compound 13.

Mass spectrometry:

<u>Compound no</u>	<u>M⁺ at m/e</u>
16	462
17	448
18	454

2-Trifluoromethanesulfonyloxy-11-(4-methylpiperazino)-5H-dibenzo[b,f][1,4]oxazepine (19) (Scheme 4). 2-Methoxy-5,10-dihydro-11-oxo-dibenzo[b,f][1,4]oxazepine (RN [60287-33-4]) was prepared according to the literature.³⁵

2-Hydroxy-5,10-dihydro-11-oxo-dibenzo[b,f][1,4]oxazepine (RN [60287-08-3]). 2-Methoxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]oxazepine (1.2 g) in 10 mL of EtSH and 10 mL of dichloromethane was treated with AlCl₃ (5.0 g) with stirring at room temperature for 6 h. The solvent of the mixture was evaporated by rotavapor at reduced pressure, quenched with 20 mL of ice-water and then 20 mL of 4N aq HCl. A white precipitate was collected, washed with water and dried under vacuum. The product (1.0 gram) showed the expected spectra including IR, ¹H NMR, and MS (EI, m/z 227).

2-Trifluoromethanesulfonyloxy-5,10-dihydro-11-oxo-dibenzo[b,f][1,4]-oxazepine. 2-Hydroxy-5,10-dihydro-11-oxo-dibenzo[b,e][1,4]oxazepine (1.0 g) in 20 mL of dichloromethane and 5 mL of triethylamine was cooled to -78 °C, treated with triflic anhydride (1.54 mL, 7 mmol). The mixture was stirred for 3 h to allow the temperature back to room temperature, quenched with 3 mL of 50% methanol in water and concentrated. The residue was flashed on silical-gel column (hexane and ethyl acetate, 9:1 then 1:1, as eluents) to afford 1.3 g of the title compound as white crystal: MS (EI) m/z 359.

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2-Trifluoromethanesulfonyloxy-11-(4-methylpiperazino)-5H-dibenzo-[b,f][1,4]oxazepine. 2-Trifluoromethanesulfonyl-5,10-dihydro-11-oxo-dibenzo-[b,e][1,4]oxazepine (1.0 g), POCl_3 (5 mL), toluene (10 mL) and N,N-dimethyl-aniline (1.0 mL) were combined and heated to reflux for 3 h. The solvent of the mixture was evaporated under vacuum to afford 5 iminochloride intermediate, which was dissolved in 10 mL of toluene and was treated with 5 mL of N-methylpiperazine under refluxing conditions for 3 h. After evaporation the residue was purified by flash chromatography (SiO_2 , 9:1 hexane and ethyl acetate then pure ethylacetate as eluents) and recrystallization (n-hexane) to afford 1.01 g of the title compound: mp 117-118 °C; MS(EI) m/z 441; ^1H NMR (CDCl_3) and elements (C, H, N) 10 were analyzed.

2-Trifluoromethanesulfonyloxy-11-(4-methylpiperazino)-5H-dibenzo-[b,f][1,4]thiazepine (20) (Scheme 5). 2-[(4-Methoxyphenyl)thio]benzoic acid (RN 19862-91-0) was prepared according to the literature.³⁸

2-[(4-methoxyphenyl)thio]benzoyl azide. 2-[(4-methoxyphenyl)thio]benzoic acid (14 g, 53.8 mmol) was treated with 50 mL of SOCl_2 and 1 mL of DMF at refluxing for 1 h. The mixture was evaporated under vacuum and the solid residue was dissolved into dry acetone (100 mL). The resulting solution was added dropwise to a cooled 30 % aq NaN_3 15 (60 mL) during 30 min. The suspension was stirred at 0 °C for 1 h, diluted with water (400 mL), filtered, washed well with water and dried overnight under vacuum to afford 14 g of the title compound as a fine solid. IR and GC-MS were analyzed. Mass spectra (EI) observed m/z at 257 ($\text{M}^+ - \text{N}_2$, isocyanation product under GC conditions: 150-300 °C/10 °C rate program) conditions.

2-Hydroxy-dibenzo[b,f][1,4]thiazepine-11(10H)-one. 2-[(4-methoxy-phenyl)thiobenzoyl azide (1.4 g) in 20 mL of dichlorobenzene was added to a suspension of AlCl_3 (2.7 g) in o-dichlorobenzene (50 mL) with stirring. The mixture was heated to reflux for 15 min, cooled to room temperature. CHCl_3 (150 mL) was added, extracted with 4 N aq HCl (2 x 30 50 mL). The organic layer was concentrated under vacuum and the residue was purified by flash chromatography (SiO_2 , ethyl acetate as eluent) to afford 240 mg of the title compound. Mass spectra observed m/z at 243 (M^+).

2-Triflate-dibenzo[b,f][1,4]thiazepine-11(10H)-one. 2-Hydroxy-dibenzo-[b,f][1,4]thiazepin-11(10H)-one (100 mg) in CH_2Cl_2 was cooled to -78 °C and treated with triflic anhydride (0.13 mL) in the presence of triethylamine (1.0 mL). After stirring for 2 h the mixture was

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mixed with SiO₂ (3 mL), vaporated and flash d (SiO₂ column, hexane and ethyl acetate as eluents) to afford 130 mg of the title compound. MS (EI) at m/z 375 (M⁺).

2-Trifluoromethanesulfonyloxy-11-(4-methylpiperazino)-5H-dibenzo-[b,f][1,4]thiazepine.
5 2-Trifluoromethanesulfonyloxy-5,10-dihydro-11-oxo-dibenzo-[b,e][1,4]thiazepine (80 mg), POCl₃ (1 mL), toluene (1 mL) and N,N-dimethyl-aniline (0.1 mL) were combined and heated to reflux for 3 h. The solvent of the mixture was evaporated under vacuum to afford iminochloride intermediate, which was dissolved in 1 mL of toluene and was treated with 1 mL of N-methylpiperazine under refluxing conditions for 3 h. After evaporation the residue was purified by flash chromatography (SiO₂, 9:1 hexane and ethyl acetate then pure ethylacetate as eluents) and recrystallization from hexane to afford 33 mg of the title compound: MS(EI) m/z 457; ¹H NMR (CDCl₃) and IR were analyzed.

Derivatives with the -CH₂CH₂-2-imidazolidinone and -CH₂CH₂CH₂-2-imidazolidinone substituent R₂ (see Formula I).

These derivatives are synthesized from the corresponding N-methyl-piperazine derivatives (e. g. compounds 12, 15-20) by first performing a demethylation with 1-chloroethyl chloroformate³⁹ and then an alkylation with the appropriate alkylating agent (e. g. 1-(2-chloroethyl)-2-imidazolidinone and 1-(3-chloropropyl)-2-imidazolidinone, respectively), refluxing over night in methyl isobutyl ketone (MIBK) in the presence of dry, ground K₂CO₃ (s) and a catalytic amount of KI (s).¹

Pharmacology

25 **Microdialysis experiments:** Adult male Wistar rats (Harlan Centraal Proefdier Bedrijf, The Netherlands) were used. Until implantation of the dialysis probe, the rats were housed in groups of 5-10 animals in plastic cages. The rats were kept in a room maintained under constant temperature (21 °C) and humidity (40%) on a 12:12 dark:light cycle (07:00 on 30 19:00 off) with food and water available ad libitum. The rats weighed between 250-350 g at the time of the microdialysis probe implantation. Animal procedures were conducted in accordance with guidelines published in the NIH Guide for Care and Use of Laboratory Animals and all protocols were approved by an Institutional Animal Care and Use Committee.

35 Drugs were administered s.c. in a volume of 1 mL/kg in distilled water. Sometimes a drop of acetic acid was added to improve the solubility of the test compound.

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- The microdialysis probes used in the present investigation were of a vertical, concentric design: A piece of fused silica capillary (Polymicro Technologies, Phoenix, AZ, USA) with I.D. 50 μm and O.D. 115 μm was threaded through a piece of PE-20 tubing (Clay Adams, Parsippany, NJ, USA) 0.5 cm from the end. A dialysis membrane (8 mm) (Hospal, Meyzieu, France) with an I.D. of 250 μm and an O.D. of 300 μm was slipped over the fused silica, glued into the PE-20, and closed at the end with epoxy (Devcon, Danvers, MA, USA). The fused silica was then brought into close apposition to the closed end of the dialysis membrane and fixed into position by epoxy at the junction of the silica and PE-20. The active membrane region was 2 mm in length and the region above the active area was coated with a thin layer of epoxy. After cutting the silica and PE-20 tubing 5 mm from the junction of the silica and PE-20, an inlet (a 25 G stainless steel needle that was broken at the bevel and from the plastic fixture) was glued with epoxy over the silica. The other end of the PE 20 served as the outlet.
- 15 The animals were anesthetized with chloral hydrate (400 mg/kg) and a microdialysis probe was implanted into the CPu at the following coordinates: (flat skull) AP 1.0 mm, ML + 2.5 mm relative to bregma, and -6.0 mm below the dura and/or nucleus accumbens at the following coordinates: (flat skull) A.P. 2.5 mm, ML -1.3 relative to bregma, and -7.3 mm below the dura. The probes were lowered using an electrode carrier (Model 1760, David Kopf, Tujunga, CA, USA) at a speed of approximately 400 $\mu\text{m}/\text{min}$ and secured to the skull with fast-curing dental cement on three skull screws. At the start of the experiment, 20 16 hours after surgery, artificial cerebrospinal fluid (aCSF) consisting of (mmol/L): NaCl 147, KCl 2.5, CaCl₂ 1.3, and MgCl₂ 0.9, pH 7.4 was perfused through the probe with a syringe pump (Harvard Apparatus Inc, South Natick, MA, USA). The outlet of the probes 25 were connected to the valve of an HPLC apparatus and samples were collected on-line and injected every 15 min from a 20 μL sample-loop. Dopamine, DOPAC, and 5-HIAA were quantified in the samples according to previously described methods. Before initiating any pharmacological manipulation, baseline samples were taken until there was less than 30% variation between the samples.
- 30 Behavioral experiments: Adult male Wistar rats (Harlan Centraal Proefdier Bedrijf, The Netherlands) were used. The rats were housed in groups of 12 animals in plastic cages. The rats were kept in a room maintained under constant temperature (21 °C) and humidity (40%) on a 12:12 dark:light cycle (07:00 on and 19:00 off) with food and water available ad libitum. The rats weighed between 200-250 g at the start of the experiment. 35 Animal procedures were conducted in accordance with guidelines published in the NIH Guide for Care and Use of Laboratory Animals and all protocols were approved by an

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Institutional Animal Care and Use Committee.

Catalepsy test: Rats were placed in a plastic bowl which measured 30 cm in diameter and 20 cm in height. After 10 minutes, the rats were placed on a vertical metal grid (grids 1 cm square) which measured 15 x 40 cm (width x height). The time until the first paw movement and the time to get off the grid were measured in three consecutive sessions. Thereafter, the rat was injected with either saline or test compound and left in the cage for another 30 minutes before catalepsy was retested.

- 10 *Locomotor activity:* Directly after the second catalepsy test (see above), the animals were transferred to a locomotion chamber (Columbus Instruments, Ohio, USA) and locomotion was measured for 10 minutes (referred to as exploratory behavior due to the new environment). Apomorphine (1 mg/kg, s.c.) was subsequently injected and locomotion was measured for 30 minutes after the injection.

15

Results

TABLE 1. Summary of the pharmacological characterization of compound 12, 15, 19, 20, clozapine, olanzapine, and haloperidol.

5

CB2	DOSE (umol/kg)	DA (1A Increase)	CATAPLEXY (car/total)	NOVELTY (count)	AMPHETAMINE		
					1 mg/kg	0.25 mg/kg	0.5 mg/kg (counts)
saline	0	0/4	439	2311	2285	1756	1752
HAL	0.03	80	1/4	416	2019	1644	2013
	0.1	115	n.d.	n.d.	1670	1100	223
	0.3	125	3/4	83	219	415	223
	1	145	n.d.	n.d.	316	165	141
	3	150	4/4	8	147	n.d.	n.d.
OLAN	0.01	31	n.d.	n.d.	n.d.	n.d.	n.d.
	0.03	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
	0.1	80	n.d.	n.d.	1815	2298	4282
	0.3	n.d.	n.d.	n.d.	1407	1768	1967
	1	85	0/4	114	323	2225	1880
	3	n.d.	n.d.	n.d.	1250	1080	2225
	10	126	0/4	9	47	491	257
	30	n.d.	n.d.	n.d.	n.d.	46	62
	100	n.d.	0/4	4	4	n.d.	n.d.
CLOZ	10	0	0/4	443	2503	2213	3503
	30	35	0/4	n.d.	n.d.	1662	1648
	100	53	1/4	46	169	1172	751
	12	0.01	12	n.d.	n.d.	n.d.	n.d.
	1	30	n.d.	n.d.	1666	1648	3587
	1	94	n.d.	n.d.	n.d.	1017	2573
	3	n.d.	n.d.	n.d.	n.d.	1403	1403
	10	144	0/4	151	2618	1284	919
	30	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
	100	n.d.	0/4	60	571	n.d.	n.d.
	100	42	No further characterization of 15, 19, and 20.				
	15	10	119	60			
	100	100					
	19	1					
	20	10					

Footnotes Table 1: Interaction with MK801: Rats were injected with test compounds 15 min prior to the administration of 0.25 mg/kg MK801 in their home cages. Immediately after the administration of MK801, the rats were transferred to a locomotor chamber (Columbus Instruments, Ohio, USA) and locomotion was measured for 30 min.

Interaction with AMPHETAMINE: Rats were injected with these compounds 15 min prior to the administration of 0.5 or 2 mg/kg amphetamine in their home cages. Immediately after the administration of amphetamine, the rats were transferred to a locomotor chamber (Columbus Instruments, Ohio, USA) and locomotion was measured for 60 min.

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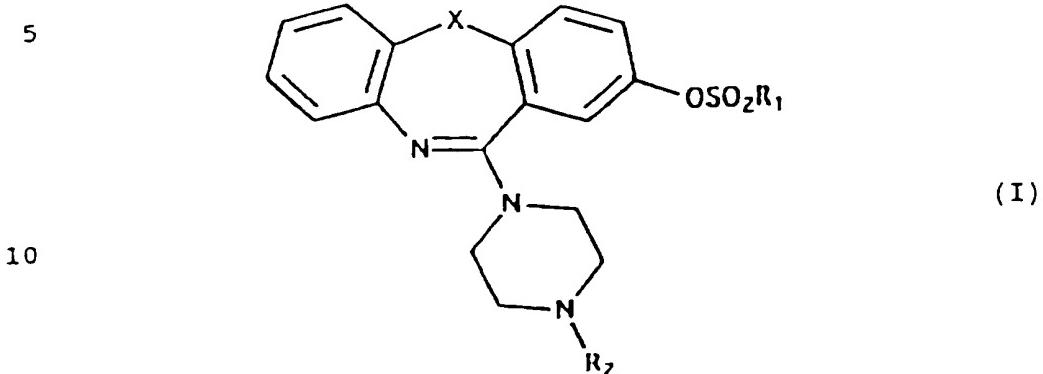
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C L A I M S

1. Compounds of Formula (1):



15 or pharmaceutically acceptable acid addition salts thereof,
wherein R₁ is H, (C₁-C₈) alkyl, haloalkyl, hydroxyalkyl,
alkenyl, alkynyl, cyclopropylalkyl, aryl, arylalkyl, hetero-
aryl, heteroarylalkyl; R₂ is H, (C₁-C₈) alkyl, alkenyl,
alkynyl, cyclopropylalkyl or (C₁-C₈) haloalkyl, hydroxyalkyl,
hydroxyalkyloxyalkyl or 1-(alkyl-2-imidazolidinone); X is NH,
20 NR₁, O, S, SO, SO₂.

2. A compound according to claim 1, wherein R₁ is selected from (C₁-C₃) alkyl and haloalkyl, R₂ is H or CH₃ and X is NH, O or S.

3. A compound according to claim 2, wherein R₁ is ClH₃ or CF₃.

30 4. A compound according to claim 1, which is selected from
2-trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-
dibenzo[b,e][1,4]diazepine,
35 2-methanesulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo
[b,e][1,4]diazepine,

2-trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazino)-
dibenzo[b,f][1,4]oxazepine,

5 2-methanesulfonyloxy-11-(4-methyl-1-piperazinyl)-dibenzo
[b,e][1,4]oxazepine,

2-trifluoromethanesulfonyloxy-11-(4-methyl-1-piperazino)-
dibenzo[b,f][1,4]thiazepine,

10 2-methanesulfonyloxy-11-(4-methyl-1-piperazino)-dibenzo-
[b,e][1,4]thiazepine,

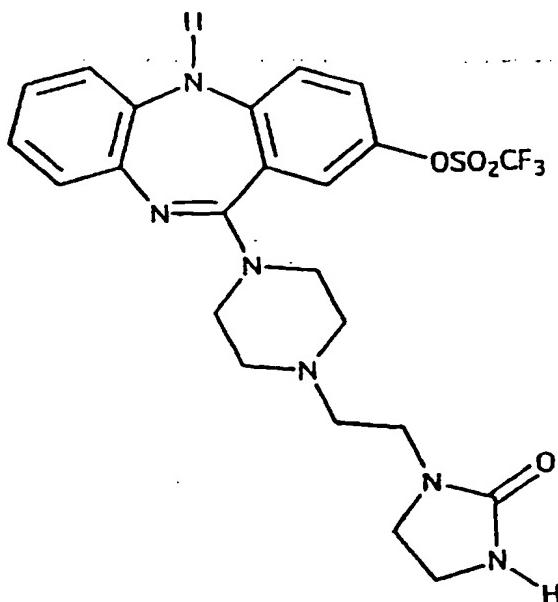
2-(4-methylphenylsulfonyloxy)-11-(4-methyl-1-piperazinyl)-
dibenzo[b,e][1,4]diazepine,

15 2-phenylsulfonyloxy-11-(4-methyl-1-piperazinyl)-5H-dibenzo-
[b,e][1,4]diazepine,

20 2-(2-thienylsulfonyloxy)-11-(4-methyl-1-piperazinyl)-5H-
dibenzo[b,e][1,4]diazepine and

the compound having the formula

25

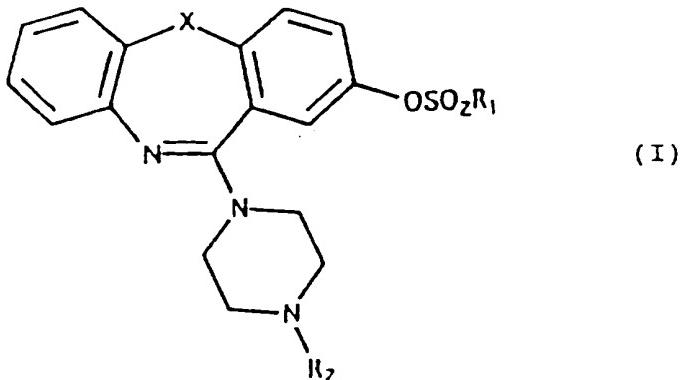


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5. Process for the preparation of a compound having the general formula (I) or a pharmaceutically acceptable acid addition salt thereof

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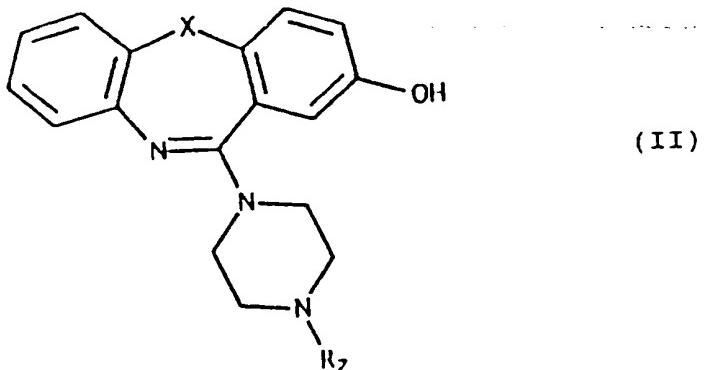


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15 wherein R₁ is H, (C₁-C₈) alkyl, haloalkyl, hydroxyalkyl, alkenyl, alkynyl, cyclopropylalkyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl; R₂ is H, (C₁-C₈) alkyl, alkenyl, alkynyl, cyclopropylalkyl, (C₁-C₈) haloalkyl, hydroxyalkyl, hydroxyalkyloxyalkyl or 1-alkyl-2-imidazolidinone; X is NH, NR₁, O, S, SO, SO₂, wherein R₁ is as defined above, which process comprises

20 a) reacting a compound of the general formula (II)

25



30

with
a1) a compound of the general formula (III)

35

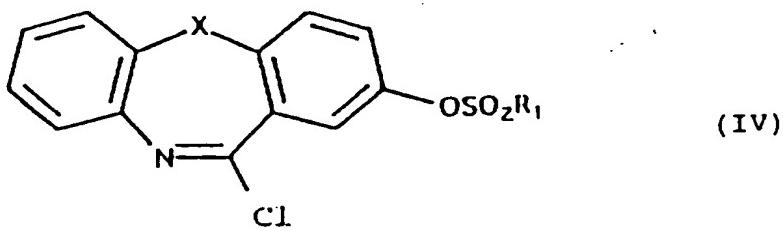


wherein R₁ is as defined above, to the formation of a compound of the general formula (I) above; or with

- 5 a2) N-phenyltrifluoromethanesulphonimide to the formation of a compound of formula (I), wherein R₁ is trifluoromethyl; or

- b) reacting a compound of the general formula (IV)

10



15

wherein R₁ and X are as defined above, with a compound of the general formula (V)

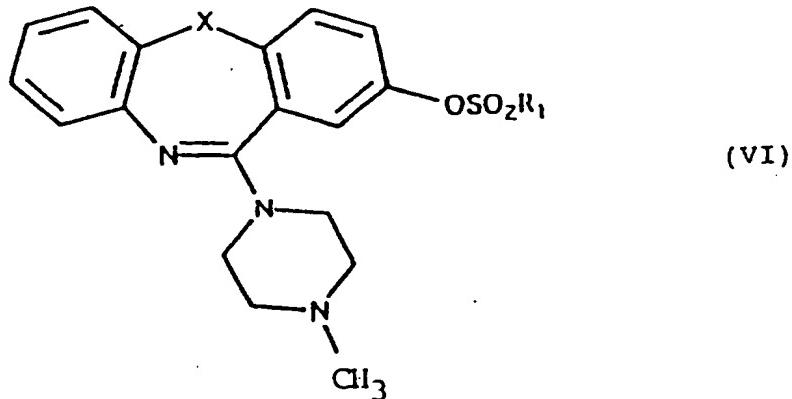
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25

- wherein R₂ is as defined above, to the formation of a compound of the general formula (I); or
- c) demethylating a compound of the general formula (VI)

30



35

wherein R₁ and X are as defined above, using 1-chloroethyl chloroformate and then alkylating with a compound of formula (VIII)

5

Y - R₂

wherein R₂ is as defined above and Y is leaving group to the formation of a compound of formula (I), wherein R₁, R₂ and X are as defined above;

10

and if, desired, converting a compound of formula (I) into another compound of formula (I), and/or, if desired, converting a compound of formula (I) into a pharmaceutically acceptable salt thereof and/or, if desired, converting a salt 15 into a free compound, and/or if desired, resolving a mixture of isomers of compounds of formula (I) into the single isomers.

6. A pharmaceutical composition containing a suitable 20 carrier and/or diluent and, as an active principle, a compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt thereof.

7. A compound of formula (I) or a therapeutically acceptable 25 salt thereof as defined in claim 1 for use in therapy.

8. Use of a compound of formula (I) or a therapeutically acceptable salt thereof as defined in claim 1, in the manufacture of medicament for use in treating psychotic/ 30 schizophrenic disorders in one or several receptor systems of the central nervous system.

9. A method for treating psychotic/schizophrenic disorders in one or several receptor systems of the central nervous 35 system in mammals, including man, which method comprises administering to the mammal of a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof as defined in claim 1.

10. The method according to claim 9, wherein said compound of formula (I) or the pharmaceutically acceptable salt thereof is administered in an amount of from about 0,1 to about 2000 mg oral daily dose, or from 0,01 to about 100 mg parenteral daily dose.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00344

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: C07D 243/38, C07D 267/20, C07D 281/16, C07D 403/12 A61K 31/495
 According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE, DK, FI, NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS-ONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	CH 422793 A (DR. A. WANDER AG), 29 April 1967 (29.04.67) --	1-8
A	GB 1216523 A (DR. A. WANDER S.A. ET AL), 23 December 1970 (23.12.70) --	1-8
A	Chemical Abstracts, Volume 93, No 17, 27 October 1980 (27.10.80), (Columbus, Ohio, USA), Protiva, Miroslav et al, "Procataleptogenic 5H-dibenzo(b,e)-1,4-diazepine derivative", page 685, THE ABSTRACT No 168316b, Czech. 1979, 179 (793) --	1-8

 Further documents are listed in the continuation of Box C. See patent family annex.

- * Special categories of cited documents
- "A" document defining the general state of the art which is not considered to be of particular relevance
- "B" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed
- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search

26 June 1996

Date of mailing of the international search report

12 -07- 1996

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00344

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>Collection Czechoslov. Chem. Commun., Volume 44, 1979, Karel Sindelár et al, "Potential Metabolites of the Neuroleptic Agents Noroxyclothepin and Oxyclothepin; Synthesis of 8-CHLORO-3-HYDROXY-10-(4-(2-HYDROXYETHYL)PIPERAZINO)- AND -10-(4-(3-HYDROXYPROPYL)PIPERAZINO)-10, 11-DIHYDRODIBENZO.....", page 3614 - page 3626, see compound XII</p> <p>--</p>	1-8
A	<p>WO 9500478 A1 (H. LUNDBECK A/S), 5 January 1995 (05.01.95), see page 12 formula U</p> <p>--</p> <p>-----</p>	1-8

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00344

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 9-10
because they relate to subject matter not required to be searched by this Authority, namely:
A method for treatment of the human or animal body by therapy, see rule 39.1.
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/SE 96/00344

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
CH-A- 422793	29/04/67	NONE		
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GB-A- 1216523	23/12/70	AT-A- 292707 AT-A- 292716 AT-A- 292717 AT-A- 292718 BE-A- 712114 CH-A- 499539 CH-A- 514612 CH-A- 517759 DE-A- 1720007 NL-A- 6406089 NL-A- 6803570 SE-B- 364277 US-A- 3546226 US-A- 3683034 US-A- 3852446 CH-A- 481133 CH-A- 481940 CH-A- 481941 CH-A- 481942 US-A- 3539573 US-A- 3751415 US-A- 3758479 US-A- 3793325 US-A- 3908010 CH-A- 484924 CH-A- 484926 CH-A- 484927 CH-A- 484928 CH-A- 484937 AT-A- 292719 AT-A- 292722 CH-A- 485749 CH-A- 485752 AT-A- 292720 AT-A- 292721 US-A- 3884920	15/08/71 15/08/71 15/08/71 15/08/71 13/09/68 30/11/70 31/10/71 15/01/72 19/05/71 01/12/64 16/09/68 18/02/74 08/12/70 08/08/72 03/12/74 15/11/69 30/11/69 30/11/69 30/11/69 10/11/70 07/08/73 11/09/73 19/02/74 23/09/75 31/01/70 31/01/70 31/01/70 31/01/70 15/08/71 15/08/71 15/02/70 15/02/70 15/08/71 15/08/71 20/05/75	-----
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WO-A1- 9500478	05/01/95	NONE		
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